USEPA REGION 4 QUALITY ASSURANCE SECTION QAPP SUPERFUND DIVISON FINAL CHECKLIST 2007

QAPP Title: Quarry Assurance Project Location: Mc Cormick, S C Originating Organization: Black & Vla- QAPP Date: Jan 2011 Receipt Date: 11 Review Date: March 2011 Reviewer: C. Jankson EPA Regional Project Manager: C. Jank EPA Project Officer: C. Jankson	LE Project PLAN BARITE HILL RIFS (1) ATRIH Specian Projects Corp.	Zev.
Topic covered in accordance with	th requirements: Yes Do	
as specified in this checklist.	d in the QAPP does not provide sufficient detail to y missing from the document.	
Element	Meets Requirements ■ Yes □ No	
A-1. Title and Approval Page	ru-Yes □ No	
Title of QAPP	EYes □ No	
Organization's Name: Both the name of the organization preparing the QAPP and the organization conducting the project or the grantee's name.	r Yes □ No	
Dated Signature of Project Manager: Both the originating organization's PM and EPA's corresponding PM and/or PO.	e EYes □ No	
Date and Signature of Quality Assurance Manager's approval for the originating entity and for EPA.	∀Yes □ No	

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Other Signatures as Needed:	Yes	□ No	·	
A-2. Table of Contents: Including Tables, Figures and Appendices	¥Yes	□ No		
A-3. Distribution List: Including Addresses of all entities or agencies requiring copies of the QAPP	⊡Yes	□ No		
A-4. Project - Task Organization				
Identifies key project personnel, specifies technical disciplines, details their roles/responsibilities and details the chain of command	Yes	□ No		
Organization chart provided: Depicts lines of authority, independence (of QA manager), and reporting responsibilities. Org- chart also contains entries for all agencies, contractors and individuals responsible for performing QAPP preparation, sample collection, laboratory analysis, data verification, review and validation, data quality assessment; and project oversight responsibilities.	Yes	□ No		,
A-5. Problem Definition/Background.				
Clearly states the particular environmental problem to be solved, decision to be made, or outcome to be achieved. Include sufficient background information to provide a historical, scientific, and regulatory perspective for this particular project.	Yes	□ No		
Provides historical and background information concerning prior environmental investigations or assessments performed at the site. Discusses the data collected from these prior investigations and identifies any additional information that may be contained in computer databases (secondary data), etc.	⊡Yes	□ No .		÷

A-6 Project/Task Description	
Provides a summary of all work to be performed, products to be produced, and the schedule for implementation. Lists the actual measurements to be made: Including in-situ field measurements, fixed laboratory measurements, or any other type of information collected as part of the project.	Yes Do
Cites applicable regulatory standards or criteria such as action limits, ARARs, PRGs, MCLs, risk assessment screening levels, etc. Must provide the actual numerical criteria for the above items.	res □ No
Identifies all instruments/equipment needed to conduct project and identifies all key study personnel (field technicians, chemists, risk assessors, engineers, project managers, quality assurance managers, etc.)	r Yes □ No
Provides work schedule for all tasks including report preparation, response to comments, etc.	Yes Do
Identifies all required reports, records, data reports, quality assurance reports/documents	reyes □ No
A-7. Data and Field Quality Objectives and Criteria for All On-Site and Off-Site Measurement Data	
Provides the Data Quality Objectives in accordance and compliance with EPA's Data Quality Objective Process (EPA-QA/G-4) document. Lists the seven steps of the DQO process and provides the project-specific information pertaining to each of these steps.	res □ No
Applies the DQO process to the project study undertaken. Provides the qualitative and quantitative data quality objectives for all aspects of the project. Must provide clearly	¥es □ No

Superfund Division QAPP Final Checklist - 2007 delineated project objectives such as determining the presence/absence of potential contaminants, nature and extent of contamination, determining whether human health is affected. Must provide a list of decisions and alternative actions (remediation, removal, further assessments, no further action, etc.). TO Yes □ No Provides all regulatory standards/criteria as part of DQO process (action limits, ARARs, PRGs, MCLs, etc.) on an analyte by analyte basis. Provides a list of all the critical ✓ Yes □ No contaminants/analytes along with their respective detection limit requirements (for chemical parameters) and QA/QC requirements. A-8. Special Training Requirements and **Special Certifications** Yes □ No Identifies how training needs are determined and lists all training requirements for the project. Specifies whether certain professionals require a license or certification to perform duties as required by federal or state laws. Yes Identifies where training records will be □ No maintained **⊡**Yes Identifies how any new training requirements □ No are communicated to program/upper management Discusses the importance of QA training and Yes □ No discusses how this training is provided.

A-9. Documentation and Records	r Yes □ No
Provides a comprehensive list of the documents and records required for this project (including raw data, field logs, audit reports, QA reports, progress or status reports, analytical data reports, data validation reports/data quality assessments reports.)	□ Yes □ No
Specifies the turnaround time for laboratory data deliverables (both hardcopy and electronic formats). Provides hardcopy data package content requirements and electronic data requirements	□ Yes □ No
Provides the retention time and location of study records, reports and formal documents.	□ Yes □ No
B-1. Sampling Process Design	
Provides a table with type and number of samples required for collection such as surface, subsurface, or groundwater.	□ Yes .□ No
Provides design of the sampling/collection network	□ Yes □ No
Provides maps or diagrams with sample locations/collection locations and provides table with frequency of sampling events	□ Yes □ No
Provides the sample matrices slated for collection in the sample table (surface soil, subsurface soil, sediment, surface water, groundwater samples, etc).	□ Yes □ No
Provides an extensive discussion regarding the rationale for the sampling design. (This also includes a discussion regarding the rationale and relevance of the analytical program).	□ Yes □ No
Provides a table identifying the chemical	□ Yes □ No

Superfund Division QAPP Final Checklist - 2007 parameters/analytes of interest for each collected sample along with the required detection limits, regulatory standards/criteria, QA/QC criteria, analytical method number, sample container requirements, sample preservation requirements, sample volume requirements and holding time criteria. **B-2. Sampling Method Requirements** ' Yes Provides the required field sample collection □ No procedures, protocols and methods ☑ Yes Provides a list of sampling/collection □ No equipment (including make and model of equipment). ☐ Yes Identifies on-site support facilities that are □ No available to field staff. T Yes Identifies key study personnel in charge of or □ No overseeing sampling/collection activities ✓ Yes Describes equipment decontamination □ No procedures and requirements. Discusses whether sampling equipment is dedicated or non-dedicated. ✓ Yes Provides table listing sample container □ No requirements and preparation requirements for these containers (if provided by laboratory, clearly states such). Yes Provides table listing sample preservation □ No requirements (for chemical parameters) and holding time criteria (where applicable). **B-3. Sample Handling and Custody** Requirements □ Yes Provides a detail description of the □ No procedures for post sample handling (once the sample has been collected). Yes Provides a detailed description of the chain-□ No of-custody procedures.

B-4. Analytical Method Requirements	
Clearly identifies the extraction, digestion, analytical methodologies (provides the actual method numbers) to be followed (includes all relevant options or modifications required), identifies the required instrumentation. Provides laboratory SOPs or QAM.	r Yes □ No
Provides validation criteria for non-standard or unpublished methodologies proposed for use for a given study.	Yes No
Identifies individual(s) responsible for overseeing the success of the analysis and for implementing corrective actions if deemed necessary.	□Yes □ No
Specifies the turnaround time for hardcopy and electronic laboratory data deliverables.	the Yes □ No
B-5. Quality Control Requirements	
Identifies the type, number and frequency of procedures and frequency of QA/QC sample collection along with the required QC statistically derived limits for each analyte (for spike samples, internal standards, surrogate spikes).	□Yes □ No
Provides the statistical equations for accuracy, precision, and comparability. Specifies the acceptance criteria for these measurements.	Yes □ No
B-6. Instrument or Equipment Testing and Inspection Requirements	∀Yes □ No
Provides a list of all in-situ testing instruments and field equipment.	⊌Yes □ No
Provides the technical criteria by which the field instruments or sampling equipment is	Yes □ No

checked for acceptable performance.			<u>.</u>	
Provides a comprehensive list of the supplies required for the project	Yes	□ No		·
Identifies the individual(s) responsible for checking and inspecting consumables and supplies	Yes	□ No		
Provides the acceptance criteria consumable item, instrument and equipment	≱ Yes	□ No		
Describes equipment and corrective maintenance practices to ensure that on-site equipment or instruments are performing within the required specifications	Yes	□ No		
Identifies the availability and location of spare parts	□ Yes	□ No		
B-7. Instrument Calibration and Frequency				
Identifies all equipment requiring calibration and discusses the frequency of calibration	Yes	□ No		
Identifies the calibration requirements for each instrument requiring calibration. (For fixed laboratory this may be in the SOPs or QA manual).	Yes	□ No		
Provides the calibration requirements and calibration acceptance criteria for each type of equipment or instrument. (Again for the off-site laboratory this information will reside in the method-specific SOPs and the QA manual).	Yes	□ No		
Identifies the type of documentation required for calibrations and instrument checks and discusses how calibrations are traced back to specific instruments for each analytical parameter. (Once again for the off-site laboratory this information will reside in the method-specific SOPs and the QA manual).	Yes	□ No		

B-8 Inspection/Acceptance Criteria and Requirements for Supplies and Consumables	□ Yes	□ No	
Provides a comprehensive list of the consumables such as, solvents, reagents, buffer solutions and other consumables or supplies required for the project.	¹ Yes	□ No	
Provides the acceptance criteria for each of these items.	Yes	□ No	·
Identifies those individual(s) responsible for checking/inspecting supplies and consumables.	₽Yes	□ No	
B-9. Data Acquisition Requirements for Non-Direct Measurements			
Identifies the type and frequency of non- direct measurement techniques for the project (for computer databases, literature searches, etc.)	₹Yes	□ No	
Clearly identified and describes the limitations of such data	ĕYes	□ No	
Discusses the rationale for using this data and explains its relevance to the project	⊉ Yes	□ No	
Specifies how limitations in this data will be communicated to all end data users and stakeholders.	₽Yes	□ No	
B-10. Data Management			
Describes the record-keeping, archival and retrieval requirements for hard-copy and electronic information produced during the course of the project.	Yes	□ No	
Provides audit checklists or other standardized forms in an appendix to the QAPP.	Yes	□ No	

Describes data handling equipment and procedures used to process, compile and analyze data (provides a complete list of computer hardware and software needs) - Specifies whether computer databases will have restricted access or will be password protected Discusses how the accuracy of computer databases is assured.	⊠ Yes	□ No
Describes process for assuring that applicable Office of Information Resource Management requirements are satisfied (mainly this is required if the data will be entered into an EPA or other Federal Database)	Yes	□ No
C-1. Assessments, Audits and Corrective Actions		
Lists the required number, frequency and type of assessments with approximate dates and names of individual(s) responsible for performing these assessments	□ Yes	□ No
Discusses one or more of the following types of assessments: peer reviews, technical audits, surveillance, management system reviews, readiness reviews, quality system audits, performance evaluations, data quality assessments.	□ Yes	□ No
Identifies the individual(s) performing these assessments and discusses the authority and independence of these individual(s) in relation to those being assessed	□ Yes	□ No
Provides a description of the types of corrective actions that may be instituted to resolve any issues raised during the audit	□ Yes	□ No
Discusses where audit findings will be documented and how the audit findings will be communicated to all key project staff, state and EPA personnel responsible for the study	□ Yes	□ No

oversight				
C-2. Reports to Management: Identifies the frequency and distribution of the following types of reports:			•	
Project Status Reports	□ Yes	□ No		
Results of Assessments or Audits	□ Yes .	□ No		
Results of periodic Data Quality Assessments	□ Yes	□ No		
QA Audit Reports	□ Yes	□ No	· · · · · · · · · · · · · · · · · · ·	
Identifies the individual(s) responsible for preparing, reviewing and receiving these reports - discusses the retention time for maintaining such reports	□ Yes	□ No		
D-1 & D-2. Data Review, Verification and Validation				
Identifies the guidance documents or SOPs governing the data review, verification and validation processes	≱ Yes	□ No		
Clearly discusses the criteria by which data will be accepted or rejected and provides a comprehensive list of the data flags or qualifiers that will be assigned to non-compliant data points (including the definitions for each of these flags)	Yes	□ No		
Describes the process, and provides the criteria by which the data will be assessed for its overall usability and intended purpose.	ĕYes	□ No		
Identifies the individual(s) responsible for validating the data and identifies the company or consultant for whom they work (Note: EPA recommends using an independent second or third party validator or at least a person that is unaffiliated with the laboratory performing the analyses on site samples).	Yes	□ No		

Identifies how problems associated with the laboratory will be documented and communicated to all end data users and stakeholders (where will the results of the data validation process be documented)	Yes	□ No	
D-3. Reconciliation of the Data to the Project-Specific DQOs	arter a		
Describes the process by which the on-site and off-site analytical data will be reconciled to the project-specific DQOs (especially if the data is non-compliant)	∠ Yes	□No	
Discusses how limitations in the final data set will be documented and communicated to all end data users and stakeholders.	∀ Yes	□ No	
Describes the circumstances under which data would be rejected and removed from the final data set	✓Yes	□ No	Tanks
Identifies the individual(s) responsible for reconciling the data to the project-specific DQOs	₽Yes	□ No	
Identifies the SOP or guidance document outlining the DQO reconciliation process	Yes	□ No	

Note: EPA's guidance and requirements documents for the DQO process, QAPP preparation, Data Validation and Data Quality Assessments, are located at www.epa.gov/quality. These documents include:

Final QAPP Disposition:

Approved, no comments	
Signature of Designated Approval Official (DAO)	04/29/11
Signature of Section Chief of the DAO	
Not Approved, Address Comments, Submit I Approval Official	Revised QAPP to the EPA Designated

References

- 1. EPA Requirements for Quality Assurance Project Plans, EPA QA/R-5, EPA/240/B-01/002 (March 2001).
- 2. EPA <u>Guidance on Systematic Planning Using the Data Quality Objectives Process</u>, EPA QA/G-4, EPA/240/B-06/001 (February 2006).

Both documents can be accessed at the following website: www.epa.gov/quality - Select guidance from the menu options to the left of the screen.